Temperature oscillations in harmonic triangular lattice with random initial velocities

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Transition to thermal equilibrium in a two-dimensional harmonic triangular lattice with nearest neighbor interactions is investigated. Initial conditions, typical for molecular dynamics simulations, are considered. Initially, particles have uncorrelated random velocities, corresponding to initial kinetic temperature of the system, and zero displacements. These initial conditions can be realized by heating of the system by an ultrafast laser pulse. In this case, the kinetic temperature of the system oscillates. The oscillations are caused by the redistribution of energy between kinetic and potential forms. At large times, energies equilibrate and temperature tends to the equilibrium value equal to a half of the initial temperature. In our previous works, an integral exactly describing this transient thermal process has been derived. The integrand depends on the dispersion relation for the lattice. The integral contains large parameter, notably time. In the present work, we investigate large time behavior of the kinetic temperature. Simple asymptotic expression for deviation of temperature from the steady state value is derived. The expression contains three harmonics with different frequencies and amplitudes. Group velocities corresponding to these frequencies are equal to zero. Two frequencies are close and therefore beats of kinetic temperature are observed. Amplitude of deviation of temperature from the steady state value decreases inversely proportional to time. It is shown that the asymptotic formula has reasonable accuracy even at small times of order of one period of atomic vibrations.

Keywords: triangular lattice, harmonic crystal, kinetic temperature, beats, transition to equilibrium, nonequilibrium process.

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Осцилляции температуры в гармонической треугольной решетке со случайными начальными скоростями

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В работе исследуется переход к состоянию теплового равновесия в двумерной бесконечной гармонической треугольной решетке с взаимодействиями ближайших соседей. Рассматриваются начальные условия, типичные для молекулярно-динамического моделирования. В начальный момент времени частицы имеют некоррелированные случайные скорости, соответствующие начальной кинетической температуре системы, и нулевые перемещения. Данные начальные условия можно рассматривать как результат воздействия на систему ультракороткого лазерного импульса. При переходе к тепловому равновесию температура совершает затухающие колебания, вызванные уравниванием кинетической и потенциальной энергий. С течением времени температура стремится к равновесному значению, равному половине начальной температуры. В предыдущих работах авторов с использованием дискретного преобразования Фурье получен интеграл, в точности описывающий данный переходный процесс. Подинтегральное выражение зависит от дисперсионного соотношения для рассматриваемой решетки. Интеграл содержит большой параметр — время, прошедшее с момента возмущения. В настоящей работе исследуется поведение температуры при больших временах. С использованием асимптотических методов получается простое выражение для отклонения температуры от равновесного значения. Выражение представляет собой сумму трех гармоник с различными частотами и амплитудами. Показано, что групповые скорости, соответствующие данным частотам, равны нулю. Две частоты являются близкими, что приводит к биениям. Амплитуда отклонения температуры от равновесного значения убывает обратно пропорционально времени. Проведено сравнение полученной асимптотической формулы с точным решением. Показано, что формула имеет приемлемую точность даже на малых временах порядка одного периода колебаний атомов.

Ключевые слова: треугольная решетка, гармонический кристалл, кинетическая температура, биения, переход к тепловому равновесию, неравновесный процесс.
1. Introduction

Analytical description of nonequilibrium thermal processes in crystals is a long-standing problem in mechanics and physics of solids [1,2]. An example of such process is a transition of a system from nonequilibrium state towards thermal equilibrium. An initial nonequilibrium state can be caused, for example, by an ultra short laser pulse [3–7] or by propagation of shock waves [8,9]. In these cases, energy is unequally distributed among degrees of freedom and the kinetic temperature can demonstrate tensor properties [8,9]. After the end of nonequilibrium process, material relaxes towards thermal equilibrium. Kinetic and potential energies tend to equilibrium values.

Similar phenomenon is observed in the beginning of molecular dynamics simulations at finite temperatures [10]. In a typical simulation, particles have random initial velocities and they are located at equilibrium positions. Then initially, total and kinetic energies of the system are equal. In the beginning of simulation, kinetic and potential energies oscillate in time and tend to the equilibrium values. The kinetic temperature, proportional to kinetic energy, also oscillates. In the present paper, we investigate this strongly nonequilibrium process. Note that the process does not depend on the initial distribution function for velocities. Behavior of the distribution function and its convergence to normal distribution is discussed in papers [11,12].

In harmonic crystals, transition to thermal equilibrium can be described analytically. In papers [13,14], the transition was investigated in harmonic one-dimensional chain. It has been shown that oscillations of kinetic temperature in a chain with random initial velocities and zero initial displacements is described by the Bessel function. In paper [15], similar problem was solved for one-dimensional chain with harmonic on-site potential. In papers [16,17] the results were generalized for the multidimensional case. In particular, an exact formula describing oscillations of kinetic temperature in harmonic triangular lattice has been derived.

In the present paper, we continue the analysis started in papers [16,17]. In-plane motions of harmonic triangular lattice with nearest-neighbor interactions are considered. Initially particles have uncorrelated random velocities, corresponding to initial kinetic temperature of the system, and zero displacements. An integral exactly describing oscillation of temperature in this system has been derived in paper [16]. In the present paper, we consider long-time asymptotic behavior of this integral. It allows, in particular, to calculate characteristic frequencies of temperature oscillations and to explain physical meaning of these frequencies.

2. Oscillations of kinetic temperature: an exact solution

In the present section, we recall the derivation of the exact expression describing oscillations of kinetic temperature in a uniformly heated harmonic triangular lattice [16].

Particles in the lattice are numbered by indices $n, k$ so that their radius vectors $r_{n,k}$ have form [18,19]

$$r_{n,k} = a(n e_1 + k e_2), \quad e_1 = e_x + e_z,$$

where $a$ is an equilibrium distance between the particles; basis unit vectors of the lattice $e_x, e_z$ are shown in Fig. 1. Each particle interacts with six nearest neighbors via harmonic forces (see Fig. 1). Anharmonicity of interactions is neglected. Anharmonic effects in triangular lattice are considered, for example, in papers [16, 20–22].

Equations of motion of the particles have form

$$\ddot{u}_{n,k} = \omega^2 (e_1 u_{n,k} - 2u_{n,k} + u_{n+1,k}) + e_2 \left( u_{n+1,k} - 2u_{n,k} + u_{n-1,k} \right) + e_1 \left( u_{n,k+1} - 2u_{n,k} + u_{n,k-1} \right),
$$

where, $\omega = \sqrt{c/m}, c$ is bond stiffness, $m$ is particle mass.

Initial velocities of the particles correspond to initial kinetic temperature of the system. Spatial distribution of temperature in the lattice is uniform (all particles have the same temperature). The velocities are independent random vectors with zero mean and equal variances. Initial displacements are equal to zero. In this case, initial kinetic energy is equal to the total energy of the system. Potential energy is equal to zero. In papers [16, 17] it is shown that the difference between kinetic and potential energies (Lagrangian) oscillates in time. It tends to zero as time tends to infinity, i.e. kinetic and potential energies equilibrate as predicted by the Virial theorem. The oscillations of the Lagrangian are described by the differential-difference equation similar to equation (1). The equation for Lagrangian is solved using the discrete Fourier transform. The exact solution allows to calculate the deviation $\delta T$ of kinetic temperature from the steady state value:

$$\delta T = T - \frac{T_n}{2} = \frac{T_n}{4\pi}(I_1 + I_2),$$

where, $T_n$ is the initial kinetic temperature of the crystal; $\tau$ is dimensionless time; $\omega$, $\omega'$ are branches of the dispersion relation for the triangular lattice defined as non-negative solutions of equation

$$\omega'^2 - 2B \omega^2 + C = 0, \quad B = \sin^2 p + \sin^2 s + \sin^2 (p + s),$$

$$C = 3 \left( \sin^2(p) + \sin^2(s) + \sin^2(p + s) \right).$$

Parameters $p, s$ are components of the wave-vector $p e_1 + s e_2$, where $e_i$ are vectors of the reciprocal basis $\mathbf{e}_i \cdot \mathbf{e}_j = \delta_{ij}$.

In the following section, we focus on the large-time asymptotic behavior of kinetic temperature. Asymptotic analysis allows to calculate the decay rate and frequencies of temperature oscillations.

Fig. 1. Particle with indices $(n,k)$ and its nearest neighbors in triangular lattice.
3. Large time behavior of kinetic temperature

In the present section, we derive simple formula describing large time behavior of the kinetic temperature using asymptotic methods [23,24].

Consider asymptotic behavior of integrals (2) at large time \((\tau \to \infty)\). We reduce the integration domain using symmetry of functions \(\omega(p, s)\). In particular, the functions satisfy the condition \(\omega(p, s) = \omega(s, p)\). Using this and other symmetry conditions, we show that integrals over 12 minimal triangles in Fig. 2 are equal. Therefore integrals \(I_\iota\) take the form

\[
I_\iota = 12 \int_{\Delta} \cos(\omega(p, s) \tau) dp ds. \tag{4}
\]

We change variables \(p, s\) to \(\omega_1, \omega_2\) in integral (4). Then the triangle \(ABC\) is mapped on the curvilinear triangle \(O_1O_2O_3\) (see Fig. 2).

Using equations (3) and (A2), we obtain the following expressions for boundaries of the integration domain (sides of the curvilinear triangle \(O_1O_2O_3\)):

\[
AB \leftrightarrow O_1O_2: \quad \omega_1 = \omega_0 \sqrt{3 - \frac{8}{9} \omega_0^2}, \quad 0 \leq \omega_0 \leq \frac{3}{2},
\]

\[
BC \leftrightarrow O_2O_3: \quad \omega_2 = \omega_0 \sqrt{3 - \frac{8}{9} \omega_0^2}, \quad \frac{3}{2} \leq \omega_0 \leq \sqrt{3}, \tag{5}
\]

\[
AC \leftrightarrow O_3O_1: \quad \omega_3 = \omega_0 \sqrt{3}, \quad 0 \leq \omega_0 \leq \sqrt{3}.
\]

To calculate Jacobian \(J\) of the transformation \(p, s\) to \(\omega_1, \omega_2\), we make two consecutive changes of variables \(p, s\) to \(B, C \) and \(B, C \) to \(\omega_1, \omega_2\). Corresponding Jacobians are calculated using formula (3)

\[
J_1 = \begin{pmatrix} p-s & p+s \\ \omega_1 & \omega_2 \end{pmatrix}^{-1}
\]

\[
J_2(\omega_1, \omega_2) = 2(\omega_1^2 + \omega_2^2) \omega_1 \omega_2.
\]

Note that \(B, C\) and \(\omega_1, \omega_2\) are related via formula (3) and Vieta’s theorem. Then the resulting Jacobian \(J_3\) is calculated as

\[
J(\omega_1, \omega_2) = \begin{vmatrix} \partial p/\partial \omega_1 & \partial s/\partial \omega_1 \\ \partial p/\partial \omega_2 & \partial s/\partial \omega_2 \end{vmatrix} = |J_1| J_2.
\]

Jacobian \(J\) is finite and it is not equal to zero inside the integration domain in formula (2). Therefore there is a unique correspondence between \(p, s\) and \(\omega_1, \omega_2\).

Changing variables in integral (4) yields

\[
I_\iota = 12 \int_{\Delta} J(\omega_1, \omega_2) \cos(\omega_1 \tau) d\omega_1 d\omega_2. \tag{6}
\]

Note that components of the inverse Jacobian are equal to components of the group velocities.

After minor algebra in formula (6), we obtain the Fourier-type integrals:

\[
I_1 = \int_{0}^{\frac{9}{4}} f_1(\omega_0) \cos(\omega_0 \tau) d\omega_0,
\]

\[
I_2 = \int_{0}^{\frac{3}{2}} f_2(\omega_0) \cos(\omega_0 \tau) d\omega_0, \tag{7}
\]

\[
f_1(\omega_0) = 12 \int_{\alpha(\omega_0)}^{\beta(\omega_0)} J(\omega_1, \omega_2) d\omega_1 d\omega_2,
\]

\[
f_2(\omega_0) = 12 \int_{\alpha(\omega_0)}^{\beta(\omega_0)} J(\omega_1, \omega_2) d\omega_1 d\omega_2,
\]

where \(\lambda_0(\omega_0), \lambda_1(\omega_0)\) are integration domains determined by formulas (5).

In the appendix, it is shown that asymptotic behavior of integrals \(I_\iota\) depends on the values of functions \(f_1\) on the boundaries of the integration domain and on singularities of these functions (see formula (A7)). It can be shown that functions \(f_\iota(\omega_0)\), defined by equations (7), are regular inside the integration domains \((0; \sqrt{3})\) and \((0; 3/2)\), except for points \(O_1, O_2, O_3\) (see Fig. 2). Points \(O_1\) and \(O_2\) do not contribute to asymptotics of integrals \(I_\iota\). The contribution is zero because \(f_1(0) = 0\) at the beginning of the integration interval and \(f_2(3/2) = 0\) at the end of the integration interval.

Functions \(f_1, f_2\) have logarithmic singularities at the points \(9/(4\sqrt{2})\) and 1 respectively. Using formulas (7), we show that in the vicinity of these points

\[
f_1 \approx -8 \sqrt{6} \ln \left| \omega_0 - \frac{9}{4\sqrt{2}} \right|, \quad f_2 \approx \frac{12}{\sqrt{7}} \ln \left| \omega_0 - 1 \right|. \tag{8}
\]

The contribution of singular points to the asymptotics is proportional to coefficients \(L_1 = 8\sqrt{6}/7, L_2 = 12/\sqrt{7}\) (see formula (A7)). Another contribution to asymptotics of the integral \(I_\iota\) is from the value of function \(f_1\) at the boundary point \(\omega = \sqrt{3}\). It is shown that

\[
\lim_{\omega \to 0} f_1(\omega_0) = 4\sqrt{3}\pi. \tag{9}
\]

Derivation of formulas (8) and (9) is given in paper [25].

Summing contributions of all special points yields:

\[
I_1 \approx 4\sqrt{3}\pi \left[ \frac{1}{\sqrt{7}} \cos \left( \frac{9}{4\sqrt{2}} \right) + \sin \left( \sqrt{3} \right) \right],
\]

\[
I_2 \approx \frac{12\pi}{\sqrt{7}} \cos \frac{\tau}{\sqrt{7}}. \tag{10}
\]

Substituting formula (10) into formula (2), we obtain the asymptotic expression describing large time behavior of the kinetic temperature:

\[
\tau \rightarrow \infty \quad \omega_0 \rightarrow \sqrt{3},
\]

\[
\phi \sim \frac{1}{\sqrt{7}} \cos \left( \frac{9}{4\sqrt{2}} \right) + \sin \left( \sqrt{3} \right),
\]

\[
\psi \approx 1, \quad \theta \approx 0.
\]

\[\tag{11}
\]

Fig. 2. Integration domain in variables \(\bar{p} = \sqrt{3} (p + s)/2, \bar{s} = (s - p)/2\) (left). Integration region in the variables \(\omega_1, \omega_2\) (right).
\[ \delta T \approx \frac{T_0}{2\sqrt{2}\pi \omega_0 t} \left[ 2\sqrt{\frac{3}{7}} \cos \left( \frac{9}{2} \omega_0 t \right) \right] + \frac{1}{3} \sqrt{3} \sin \left( 2\sqrt{6} \omega_0 t + \frac{3}{2} \right) \cos \left( 2\sqrt{3} \omega_0 t \right). \]  

Formula (11) describes the deviation of temperature from the steady-state value. It shows that the deviation decays inversely proportional to time. Oscillations of temperature have three main frequencies

\[ \Omega_1 = \frac{9}{2} \omega_0, \quad \Omega_2 = 2\sqrt{6} \omega_0, \quad \Omega_3 = 2\sqrt{3} \omega_0. \]  

These frequencies correspond to three special points of integrals \( I \). Note that frequencies \( \Omega_1 \) and \( \Omega_2 \) are close. Therefore beats of kinetic temperature are observed.

We show using formulas (3) that group velocities corresponding to frequencies (12) are equal to zero:

\[ \frac{\partial \omega_1}{\partial p} = \frac{\partial \omega_2}{\partial s}, \quad \frac{\partial \omega_1}{\partial s} = \frac{\partial \omega_2}{\partial p}, \quad \frac{\partial \omega_1}{\partial p} \neq \frac{\partial \omega_2}{\partial s} \approx 0. \]

Note that the same fact is observed in one-dimensional chain with harmonic on-site potential [15].

Comparison of the exact expression (7) for \( \delta T \) with asymptotic formula (11) is shown in Fig. 3. For \( t > 2.5t_1 \), where \( t_1 = 2\pi/\omega_0 \), the difference between exact and asymptotic formulas is less than 6%.

Thus formula (11) describes oscillations of kinetic temperature even at small times \( t \sim t_1 \).

4. Conclusions

Oscillations of kinetic temperature in harmonic triangular lattice with random initial velocities and zero initial displacements were considered. Simple asymptotic formula describing large time behavior of these oscillations was derived. Comparison with an exact solution shows that the asymptotic formula has reasonable accuracy even at small times of order of one period of atomic vibrations. It is shown that deviation of kinetic temperature from the steady state value is represented as a sum of three harmonics with different frequencies and amplitudes. The amplitudes are inversely proportional to time. Group velocities corresponding to frequencies in asymptotics are equal to zero. The same effect has previously been reported for one-dimensional chain with harmonic on-site potential [15]. Therefore we assume that in harmonic crystals frequencies of oscillations of kinetic temperature correspond to zero group velocities. However this hypothesis is awaiting a rigorous proof.

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References

Appendix. Asymptotics of Fourier-type integrals

In the present appendix, we consider asymptotic behavior of the Fourier-type integral

\[ I = \int_{0}^{b} f(\omega) \cos(\omega \tau) d\omega \]  

(A1)

for large values of parameter \( \tau \). Assume that \( f(\omega) \) is a polynomial. Then integration by parts in formula (A1) yields the following asymptotics:

\[ I = f(b) \frac{\sin b \tau}{\tau} + O\left(\frac{1}{\tau}\right), \quad \tau \to \infty. \]  

(A2)

Formula (A2) is valid for polynomial of any degree. Therefore it is also valid for function \( f(\omega) \) being regular on the interval \([0, b]\).

Formula (A2) is generalized for functions having \( N \) discontinuities on the integration interval. Assume that functions \( f \) has \( N \) finite jumps \( G_i \) at points \( \omega_i \in (0, b) \):

\[ G_i = \lim_{\varepsilon \to 0^+} \left( f(\omega + \varepsilon) - f(\omega - \varepsilon) \right). \]  

(A3)

Then separating the interval \((0, b)\) into \( N + 1 \) intervals, where \( f \) is regular, and using formula (A2) yields:

\[ I \approx f(b) \frac{\sin b \tau}{\tau} - \frac{1}{\tau} \sum_{m=1}^{N} G_i \sin \omega_i \tau. \]  

(A4)

Formula (A4) shows that asymptotic behavior of integral (A1) depends on discontinuities of function \( f \) and its value \( f(b) \) at the boundary point.

Consider logarithmic singularity of function \( f(\omega) \) at the point \( \omega = \omega', (0 \leq \omega' < b) \). In other words, in the vicinity of this point the function has form

\[ f(\omega) \approx L \ln|\omega - \omega'|. \]  

(A5)

Substituting this expression into the integral (A1) yields:

\[ I \approx \frac{L}{\tau} \left[ -\pi \cos \omega' \tau + \ln|b - \omega'| \sin b \tau, \quad \omega' > 0, \right. \]
\[ \left. -\frac{\pi}{2} + \ln|b| \sin b \tau, \quad \omega' = 0. \right] \]  

(A6)

The contribution of the logarithmic singular point to asymptotics of the integral (A1) is given by the first terms in these expressions. The second terms is the contribution of the end of the integration interval \( \omega = b \).

Generalizing formula (A4) for function \( f(\omega) \) having \( N \) discontinuities (each of them is jump or logarithmic singularity) at points \( \omega_i \in [0, b], i = 1 \ldots N \), yields:

\[ \int_{0}^{b} f(\omega) \cos(\omega \tau) d\omega \approx f(b) \frac{\sin b \tau}{\tau} - \frac{1}{\tau} \sum_{m=1}^{N} G_i \sin \omega_i \tau + \pi L \cos \omega_i \tau. \]  

(A7)

Coefficients \( G_i, L_i \) are defined by formulas (A3), (A5) respectively. If there is no logarithmic singularity at \( \omega_i \), then \( L_i = 0 \). If \( \omega_i = 0 \), then corresponding coefficient \( L_i \) in formula (A7) should be replaced by \( L_i/2 \) (see formula (A6)).